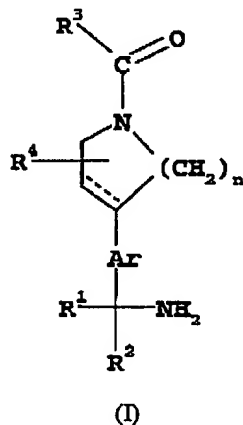
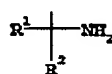


AMENDMENTS TO THE CLAIMS

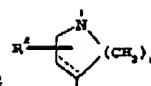
1. (Currently Amended) A compound of formula (I): $[[-]]$



such that Ar is an aryl or a heteroaryl, and the
on the aryl,
wherein $[[-]]$,



is beta to the



----- is a single or a double bond;

R¹ and R² are each independently hydrogen or lower alkyl;

R³ is aryl, arylalkenyl, cycloalkenyl, cycloalkyl, heteroaryl, heteroarylalkenyl, heterocycloalkenyl, a carbon linked heterocycloalkyl or alkyl optionally substituted by one or more groups selected from hydroxy, alkoxy, alkyloxycarbonylamino, cycloalkyl, heterocycloalkyl, R⁶, -OR⁶, -S(O)_mR⁶ or -C(=O)-R⁶;

R⁴ is hydrogen, acyl, alkoxy, alkyloxycarbonyl, carboxy, cyano, halo, hydroxy, -C(=O)-NY¹Y² or alkyl optionally substituted with alkoxy, alkylcarbonylamino, alkylsulfonylamino, hydroxy, -S(O)_m-alkyl or -NY¹Y²;

R⁶ is aryl or heteroaryl;

Y^1 and Y^2 are independently hydrogen, alkenyl, alkyl, aryl, arylalkyl, cycloalkyl, heteroaryl, heteroarylalkyl or heterocycloalkyl; or the group $-NY^1Y^2$ may form a cyclic amine; m is zero or an integer 1 to 2; and

n is 2; or

an N-oxide of said compound, a prodrug of said compound, a pharmaceutically acceptable salt of said compound, a solvate of said compound, or a hydrate of said compound.

2. (Previously Presented) The compound of Claim 1, wherein R^1 or R^2 is hydrogen, or R^1 and R^2 are hydrogen, and R^3 is an aryl or a heteroaryl.

3. (Previously Presented) The compound of Claim 2, wherein said R^3 is a phenyl or a naphthyl.

4. (Previously Presented) The compound of Claim 2, wherein said R^3 is aryl substituted with at least one substituent.

5. (Original) The compound of Claim 4, wherein said substituent is selected from the group consisting of a halo atom, an alkyl substituted by aryl, an alkyl substituted by aryloxy, an alkyl substituted by aroyl, an alkyl substituted by heteroaryl, an arylalkynyl, a heteroarylalkynyl, an aryl, a heteroaryl, an arylalkenyl and an arylalkyloxy.

6. (Original) The compound of Claim 5, wherein said aryl or heteroaryl of said substituent is further substituted by at least one aryl group substituent.

7. (Previously Presented) The compound of Claim 2, wherein said heteroaryl is a pyridyl, a quinoliny, a thienyl, a furanyl, or an indolyl.

8. (Original) The compound of Claim 7, wherein said heteroaryl is substituted with at least one substituent.

9. (Previously Presented) The compound of Claim 8, wherein said substituent is an alkyl, an alkyl substituted by an aryl, an alkyl substituted by an aryloxy, an alkyl substituted

by an aroyl, an alkyl substituted heteroaryl, an arylalkynyl, a heteroarylalkynyl, a heteroaryl, an arylalkenyl or an arylalkoxy.

10. (Original) The compound of Claim 9, wherein said aryl of said substituent is further substituted by at least one aryl substituent.

11. (Previously Presented) The compound of Claim 1, wherein R⁴ is hydrogen or a cyano group.

12. (Cancelled).

13-14. (Cancelled).

15. (Previously Presented) The compound of Claim 1, wherein:

Ar is a phenyl group;

R¹ and R² are both hydrogen;

R³ is an aryl, a naphthyl or a heteroaryl;

R⁴ is hydrogen or a cyano; and

----- is a single bond.

16. (Previously Presented) The compound of Claim 15, wherein R³ as aryl or naphthyl is substituted with at least one substituent selected from the group consisting of a halo atom, an alkyl substituted by aryl, an alkyl substituted by aryloxy, an alkyl substituted by aroyl, an alkyl substituted by aryloxy, an alkyl substituted by aroyl, an alkyl substituted by a heteroaryl, an arylalkynyl, a heteroarylalkynyl, an aryl, a heteroaryl, an arylalkenyl, and an arylalkyloxy.

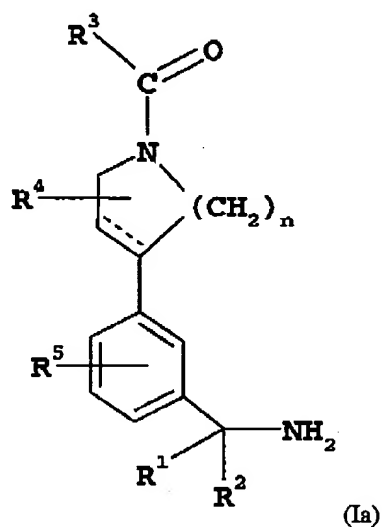
17. (Original) The compound of Claim 16, wherein said aryl or said heteroaryl of said substituent is further substituted by at least one aryl substituent.

18. (Currently Amended) The compound of Claim 15, wherein R³ as heteroaryl is substituted by at least one substituent selected from the group consisting of a pyridyl, a quinoliny, a thienyl, a furanyl, and an indolyl.

19. (Previously Presented) The compound of Claim 18, wherein said substituent of said heteroaryl is further substituted by at least one moiety selected from the group consisting of an alkyl substituted by an aryl, an alkyl substituted by an aryloxy, an alkyl substituted by an aroyl, an alkyl substituted heteroaryl, an arylalkynyl, a heteroarylalkynyl, a heteroaryl, an arylalkenyl, and an arylalkyloxy.

20. (Original) The compound of Claim 19, wherein an aryl of said moiety is further substituted by at least one aryl substituent.

21. (Currently Amended) A compound of formula (Ia):



wherein n

----- is a single or a double bond;

R^1 and R^2 are each independently hydrogen or lower alkyl;

R^3 is aryl, arylalkenyl, cycloalkenyl, cycloalkyl, heteroaryl, heteroarylalkenyl, heterocycloalkenyl, a carbon linked heterocycloalkyl or alkyl optionally substituted by one or more groups selected from hydroxy, alkoxy, alkyloxycarbonylamino, cycloalkyl, heterocycloalkyl, R^6 , $-OR^6$, $-S(O)_mR^6$ or $-C(=O)-R^6$;

R⁴ is hydrogen, acyl, alkoxy, alkyloxycarbonyl, carboxy, cyano, halo, hydroxy, -C(=O)-NY¹Y² or alkyl optionally substituted with alkoxy, alkylcarbonylamino, alkylsulfonylamino, hydroxy, -S(O)_m-alkyl or -NY¹Y²;

R⁵ is hydrogen, acyl, alkoxy, alkyloxycarbonyl, aryl, carboxy, cyano, halo, heteroaryl, heteroaryloxy, heterocycloalkyl, heterocycloalkyloxy, heterocycloalkylalkyloxy, heteroarylalkyloxy, hydroxy, trifluoromethyl, -C(=O)-NY¹Y², -NY¹Y², -Z¹-C₂₋₆alkylene-R⁷ or alkyl optionally substituted with alkoxy, alkylcarbonylamino, alkylsulfonylamino, aryl, heteroaryl, heterocycloalkyl, hydroxy, ureido, -C(=O)-NY¹Y², -SO₂-NY¹Y², -S(O)_m-alkyl or -NY¹Y², and

R⁶ is aryl or heteroaryl;

R⁷ is hydroxy, alkoxy, ureido, -C(=O)-NY¹Y², -SO₂-NY¹Y², -S(O)_m-alkyl or -NY¹Y²,

R⁸ is hydrogen or lower alkyl;

Y¹ and Y² are independently hydrogen, alkenyl, alkyl, aryl, arylalkyl, cycloalkyl, heteroaryl, heteroarylalkyl or heterocycloalkyl; or the group -NY¹Y² may form a cyclic amine;

Z¹ is O, S(O)_m or NR⁸;

m is zero or an integer 1 to 2; and

n is 2; or

an N-oxide of said compound, a prodrug of said compound, a pharmaceutically acceptable salt of said compound, a solvate of said compound, an N-oxide of said solvate of said compound, or a prodrug of said solvate of said compound.

22. (Previously Presented) The compound of Claim 21, wherein R³ is a phenyl or a naphthyl.

23. (Previously Presented) The compound of Claim 22, wherein said aryl is substituted by at least one substituent selected from the group consisting of a halo atom, an alkyl substituted by an aryl, and an alkyl substituted by a heteroaryl.

24. (Original) The compound of Claim 23, wherein said aryl or heteroaryl of said substituent is further substituted by at least one aryl group substituent.

25. (Previously Presented) The compound of Claim 21, wherein R^3 is phenylC₁₋₃alkylpyridyl, phenylC₁₋₃alkylthienyl or indolyl.

26. (Previously Presented) The compound of Claim 21, wherein R^3 is a heteroaryl selected from the group consisting of a pyridyl, a quinoliny, a thienyl, a furanyl, and an indolyl.

27. (Previously Presented) The compound of Claim 26, wherein said heteroaryl is substituted by at least one substituent selected from the group consisting of an alkyl substituted by an aryl, and an alkyl substituted by a heteroaryl.

28. (Original) The compound of Claim 27, wherein said aryl and said heteroaryl of said substituent are further substituted by at least one aryl group substituent.

29. (Previously Presented) The compound of Claim 28, wherein R^3 is phenylC₁₋₃alkylpyridyl, phenylC₁₋₃alkylthienyl or indolyl.

30. (Previously Presented) The compound of Claim 21, wherein R^4 is a hydrogen or a cyano.

31. (Previously Presented) The compound of Claim 21, wherein R^5 is a hydrogen, a lower alkyl or a halo.

32. (Previously Presented) The compound of Claim 31, wherein R^5 is methyl or fluoro.

33. (Original) The compound of Claim 31, wherein R^5 is attached to the phenyl ring of formula (Ia) in the position para to the CH₂NH₂ group.

34. (Previously Presented) The compound of Claim 21, wherein:

R^3 is a phenyl, a naphthyl, a heteroaryl selected from the group consisting of a pyridyl, a quinolinyl, a thienyl, a furanyl, and an indolyl, a phenyl substituted by at least one substituent, a naphthyl substituted by at least one substituent, or a heteroaryl selected from the group consisting of a pyridyl, a quinolinyl, a thienyl, a furanyl, and an indolyl, that is substituted by at least one substituent,

wherein said substituent is selected from the group consisting of a halo atom, an alkyl substituted by aryl, and alkyl substituted heteroaryl, wherein the aryl or heteroaryl groups are further substituted by one or more aryl group substituents;

R^4 is hydrogen or a cyano; and

R^5 is hydrogen, a lower alkyl or a halo.

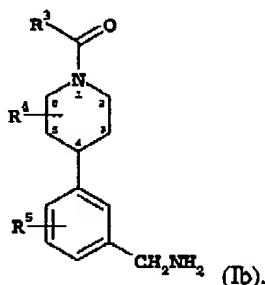
35. (Previously Presented) The compound of Claim 34, wherein:

R^3 is phenylC₁₋₃alkylpyridyl, phenylC₁₋₃alkylthienyl or indolyl;

R^4 is a hydrogen or a cyano; and

R^5 is a methyl or a fluoro, and is attached to the phenyl ring of formula (Ia) in the position para to the CR¹R²NH₂ group.

36. (Previously Presented) The compound of Claim 21, having formula (Ib):



37. (Currently Amended) The compound of Claim 36, ~~wherein R^3 as is selected from the group consisting of a phenyl and a naphthyl,~~ wherein R^3 is a phenyl or a naphthyl.

38. (Previously Presented) The compound of Claim 37, wherein said aryl is substituted by at least one substituent selected from the group consisting of a halo atom, an alkyl substituted by an aryl, and an alkyl substituted by a heteroaryl.

39. (Original) The compound of Claim 38, wherein said aryl or heteroaryl of said substituent is further substituted by at least one aryl group substituent.

40. (Previously Presented) The compound of Claim 36, wherein R^3 is a pyridyl, a quinolinyl, a thienyl, a furanyl, or an indolyl.

41. (Previously Presented) The compound of Claim 39, wherein said heteroaryl is substituted by at least one substituent selected from the group consisting of an alkyl substituted by an aryl, and an alkyl substituted by a heteroaryl.

42. (Previously Presented) The compound of Claim 39, wherein said aryl and said heteroaryl of said substituent are further substituted by at least one aryl group substituent.

43. (Previously Presented) The compound of Claim 36, wherein R^3 is phenylC₁₋₃alkylpyridyl, phenylC₁₋₃alkylthienyl or indolyl.

44. (Previously Presented) The compound of Claim 36, wherein R^4 is a hydrogen or a cyano.

45. (Previously Presented) The compound of Claim 36, wherein R^5 is a hydrogen, a lower alkyl or a halo.

46. (Previously Presented) The compound of Claim 45, wherein R^5 is a methyl or a fluoro.

47. (Original) The compound of Claim 45, wherein R^5 is attached to the phenyl ring of formula (Ib) in the position para to the CH₂NH₂ group.

48. (Previously Presented) The compound of Claim 36, wherein:

R^3 is a phenyl, a naphthyl, a heteroaryl selected from the group consisting a pyridyl, a quinolinyl, a thienyl, a furanyl, and an indolyl, a phenyl substituted by at least one substituent, a naphthyl substituted by at least one substituent, or a heteroaryl selected from the group consisting a pyridyl, a quinolinyl, a thienyl, a furanyl, and an indolyl, that is substituted by at least one substituent,

wherein said substituent is selected from the group consisting of a halo atom, an alkyl substituted by aryl, and alkyl substituted heteroaryl, wherein the aryl or heteroaryl groups are further substituted by one or more aryl group substituents;

R^4 is hydrogen or a cyano; and

R^5 is hydrogen, a lower alkyl or a halo.

49. (Previously Presented) The compound of Claim 48, wherein:

R³ is phenylC₁₋₃alkylpyridyl, phenylC₁₋₃alkylthienyl or indolyl;

R⁴ is a hydrogen or a cyano; and

R⁵ is a methyl or a fluoro.

50. (Currently Amended) The compound of Claim 1, selected from the group consisting of:

3-[1-(5-phenylethynyl-pyridine-3-carbonyl)-piperidin-4-yl]-benzylamine;

3-[1-(3-phenylethyl-benzoyl)-piperidin-4-yl]-benzylamine;

3-[1-[3-(4-hydroxyphenyl)ethyl-benzoyl]-piperidin-4-yl]-benzylamine;

3-[1-[3-(6-amino-pyridin-3-yl)ethyl-benzoyl]-piperidin-4-yl]-benzylamine;

3-[1-(5-phenylethyl-thiophene-2-carbonyl)-piperidin-4-yl]-benzylamine;

4-fluoro-3-[1-(5-phenylethyl-pyridine-3-carbonyl)-piperidin-4-yl]-benzylamine;

4-methyl-3-[1-(5-phenylethyl-pyridine-3-carbonyl)-piperidin-4-yl]-benzylamine;

3-[1-(indole-6-carbonyl)-piperidin-4-yl]-benzylamine;

4-(3-aminomethyl-phenyl)-1-(5-phenethyl-pyridine-3-carbonyl)-piperidine-4-carbonitrile;

[4-(3-aminomethylphenyl)piperidin-1-yl]-(3,4-dichlorophenyl)methanone;

1-[1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-methanoyl]-3-methylsulfanyl-6,7-dihydro-5H-benzo[c]thiophen-4-one trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(3-methylsulfanyl-6,7-dihydro-benzo[c]thiophen-1-yl)-methanone trifluoroacetate;

1-[1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-methanoyl]-3-ethylsulfanyl-6,6-dimethyl-6,7-dihydro-5H-benzo[c]thiophen-4-one trifluoroacetate;

1-{1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-methanoyl}-3-propylsulfanyl-6,7-dihydro-5H-benzo[c]thiophen-4-one trifluoroacetate;

1-{1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-methanoyl}-3-isopropylsulfanyl-6,7-dihydro-5H-benzo[c]thiophen-4-one trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-benzo[b]thiophen-2-yl-methanone-trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-4-hydroxy-piperidin-1-yl]-1-(5-phenethyl-pyridin-3-yl)-methanone-ditrifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(1-methyl-1H-indol-3-yl)-methanone-trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-[3-(2-fluoro-phenylethynyl)-phenyl]-methanone trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-[3-[2-(2-fluoro-phenyl)-ethyl]-phenyl]-methanone trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-[3-[2-(6-amino-pyridin-3-yl)-ethyl]-phenyl]-methanone tri-trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(6-chloro-thieno[3,2-*b*]thiophen-2-yl)-methanone trifluoroacetate;

(3R,4S) and (3S, 4R)-4-(3-Aminomethyl-phenyl)-1-(5-phenethyl-pyridine-3-carbonyl)-piperidine-3-carboxylic acid ethyl ester dihydrochloride;

3-[1-(5-Phenylethynyl-furan-2-carbonyl)-piperidin-4-yl]-benzylamine trifluoroacetate;

4-(3-Aminomethyl-phenyl)-piperidine-1-carboxylic acid (3,4-dichloro-phenyl)-amide trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(2,3-dihydro-benzofuran-5-yl)-methanone;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(5,6-dichloro-pyridin-3-yl)-methanone;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(3-bromo-4-fluoro-phenyl)-methanone;

(E)-1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-3-(2-nitro-phenyl)-propenone;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(3-bromo-5-iodo-phenyl)-methanone; and

(E)-1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-3-phenyl-propenone.

51. (Previously Presented) A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 1 and a pharmaceutically acceptable carrier thereof.

52. (Currently Amended) The pharmaceutical composition of Claim 51, wherein said compound is selected from the group consisting of:

3-[1-(5-phenylethynyl-pyridine-3-carbonyl)-piperidin-4-yl]-benzylamine;

3-[1-(3-phenylethyl-benzoyl)-piperidin-4-yl]-benzylamine;

3-[1-[3-(4-hydroxyphenyl)ethyl-benzoyl]-piperidin-4-yl]-benzylamine;

3-[1-[3-(6-amino-pyridin-3-yl)ethyl-benzoyl]-piperidin-4-yl]-benzylamine;

3-[1-(5-phenylethyl-thiophene-2-carbonyl)-piperidin-4-yl]-benzylamine;

4-fluoro-3-[1-(5-phenylethyl-pyridine-3-carbonyl)-piperidin-4-yl]-benzylamine;

4-methyl-3-[1-(5-phenylethyl-pyridine-3-carbonyl)-piperidin-4-yl]-benzylamine;

3-[1-(indole-6-carbonyl)-piperidin-4-yl]-benzylamine;

4-(3-aminomethyl-phenyl)-1-(5-phenethyl-pyridine-3-carbonyl)-piperidine-4-carbonitrile;

[4-(3-aminomethylphenyl)piperidin-1-yl]-(3,4-dichlorophenyl)methanone;

1-[1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-methanoyl]-3-methylsulfanyl-6,7-dihydro-5H-benzo[c]thiophen-4-one trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(3-methylsulfanyl-6,7-dihydro-benzo[c]thiophen-1-yl)-methanone trifluoroacetate;

1-[1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-methanoyl]-3-ethylsulfanyl-6,6-dimethyl-6,7-dihydro-5H-benzo[c]thiophen-4-one trifluoroacetate;

1-[1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-methanoyl]-3-propylsulfanyl-6,7-dihydro-5H-benzo[c]thiophen-4-one trifluoroacetate;

1-[1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-methanoyl]-3-isopropylsulfanyl-6,7-dihydro-5H-benzo[c]thiophen-4-one trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-benzo[b]thiophen-2-yl-methanone-trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-4-hydroxy-piperidin-1-yl]-1-(5-phenethyl-pyridin-3-yl)-methanone-ditrifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(1-methyl-1H-indol-3-yl)-methanone-trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-[3-(2-fluoro-phenylethynyl)-phenyl]-methanone trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-[3-[2-(2-fluoro-phenyl)-ethyl]-phenyl]-methanone trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-[3-[2-(6-amino-pyridin-3-yl)-ethyl]-phenyl]-methanone tri-trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(6-chloro-thieno[3,2-*b*]thiophen-2-yl)-methanone trifluoroacetate;

(3R,4S) and (3S, 4R)-4-(3-Aminomethyl-phenyl)-1-(5-phenethyl-pyridine-3-carbonyl)-piperidine-3-carboxylic acid ethyl ester dihydrochloride;

3-[1-(5-Phenylethynyl-furan-2-carbonyl)-piperidin-4-yl]-benzylamine trifluoroacetate;

4-(3-Aminomethyl-phenyl)-piperidine-1-carboxylic acid (3,4-dichloro-phenyl)-amide trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(2,3-dihydro-benzofuran-5-yl)-methanone;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(5,6-dichloro-pyridin-3-yl)-methanone;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(3-bromo-4-fluoro-phenyl)-methanone;

(E)-1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-3-(2-nitro-phenyl)-propenone;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(3-bromo-5-iodo-phenyl)-methanone;

(E)-1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-3-phenyl-propenone; and

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-3-cyclohexyl-propan-1-one.

53. (Previously Presented) A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 21 and a pharmaceutically acceptable carrier thereof.

54. (Previously Presented) A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 36 and a pharmaceutically acceptable carrier thereof.

55. (Previously Presented) A method for treating asthma in a patient comprising administering to the patient a therapeutically effective amount of a compound of Claim 1.

56-57. (Cancelled)

58. (Previously Presented) A method for treating asthma in a patient comprising administering to the patient a therapeutically effective amount of a compound of Claim 21.

59-60. (Cancelled)

61. (Previously Presented) A method for treating asthma in a patient comprising administering to the patient a therapeutically effective amount of a compound of Claim 36.

62-72. (Cancelled)

73. (Previously Presented) The compound of Claim 25, wherein the phenylC₁₋₃alkylpyridyl is 5-phenylethyl-pyrid-3-yl, the phenylC₁₋₃alkylthienyl is and the indolyl is indol-6-yl.

74. (Previously Presented) The compound of Claim 29, wherein the phenylC₁₋₃alkylpyridyl is 5-phenylethyl-pyrid-3-yl, the phenylC₁₋₃alkylthienyl is 5-phenylethyl-thien-2-yl and the indolyl is indol-6-yl.

75. (Previously Presented) The compound of Claim 35, wherein the phenylC₁₋₃alkylpyridyl is 5-phenylethyl-pyrid-3-yl, the phenylC₁₋₃alkylthienyl is 5-phenylethyl-thien-2-yl and the indolyl is indol-6-yl.

76. (Previously Presented) The compound of Claim 1, wherein ----- is a single bond.